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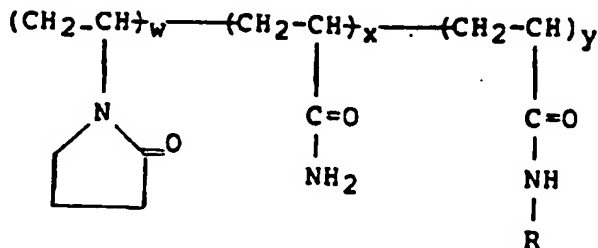
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Terpolymers.

Novel N-vinyl pyrrolidone/acrylamide/N-alkyl acrylamide terpolymers have the formula:



wherein w is 1 to 80 mole percent, preferably 10 to 70 mole percent; x is 10 to 90 mole percent, preferably 30 to 70 mole percent; and y is 0.1 to 10.0 mole percent, preferably 0.2 to 3.0 mole percent; R is alkyl groups having 6 to 30 carbon atoms, preferably 8 to 18.

The terpolymers are useful as improved viscosification agents for aqueous solutions containing, for example, an acid or a base or a salt.

The viscosification agents of the invention are normally water-soluble or water-dispersible.

$$\begin{array}{ccccc} (\text{CH}_2-\text{CH})_w & - & (\text{CH}_2-\text{CH})_x & - & (\text{CH}_2-\text{CH})_y \\ | & & | & & | \\ \text{N} & & \text{C=O} & & \text{C=O} \\ | & & | & & | \\ \text{O} & & \text{NH}_2 & & \text{NH} \\ & & & & | \\ & & & & \text{R} \end{array}$$

The polymer molecular weights, as indicated by intrinsic viscosities and confirmed by sedimentation/light scattering, for the terpolymers of acrylamide/N-vinyl pyrrolidone/N-alkyl acrylamide is 10^3 to 2×10^7 , more preferably 10^4 to 1.5×10^7 , and most preferably 10^5 to 5×10^6 . The molecular weight of the terpolymers is determined from solution viscosity measurements via standard methods and confirmed by sedimentation/light scattering. Specifically, following the initial isolation of the water soluble terpolymers they are redissolved in salt water (e.g., 2% NaCl or 3% NaCl + 0.3% CaCl₂) to give solutions with known concentrations. The low shear viscosities of the solutions and the pure solvent were measured in a standard Contraves viscometer. Subsequently, the reduced viscosity is calculated through standard methods utilizing these values. Extrapolation to zero polymer concentration leads to the intrinsic viscosity of the polymer solution. The intrinsic viscosity is directly related to the molecular weight through the well-known Mark-Houwink relationship.

The intrinsic viscosity of the terpolymers (.15% polymer, 3.0% NaCl, 0.3% CaCl_2) at 1.28s^{-1} is 5 to 20.

The terpolymers of the present invention have un-expectedly high viscosification power, for example in brine solutions.

Terpolymerization of the acrylamide monomer, N-vinyl pyrrolidone monomer, and N-alkyl acrylamide monomer is preferably effected at a temperature of 10°C to 90°C, more preferably at 10°C to 70°C, and most preferably 20°C to 60°C for a period of time of preferably 1 to 24 hours, more preferably 2 to 10 hours, and most preferably 3 to 8 hours.

5 Although NVP monomer will dissolve substantial quantities of long chain N-alkylacrylamide monomers (e.g., *n*-octylacrylamide), the addition of water and AM monomer often leads to a cloudy, nonhomogeneous reaction mixture. Addition of SDS clarifies the solution and solubilizes the hydrophobic monomer in the aqueous phase, thereby improving the process. Also, the use of SDS leads to a more gel-free product as judged by visual inspection.

10 A suitable method of recovery of a formed water soluble terpolymer from the aqueous reaction solution comprises precipitation in acetone, methanol, ethanol and the like.

Suitable water or oil soluble free radical initiators for the free radical terpolymerization of the acrylamide monomers, the N-vinyl pyrrolidone, and the N-alkyl acrylamide monomer are selected from the group consisting of azo compounds, peroxides and persulfates. However, the preferred initiators are azo compounds, such as 2,2'-azobisisobutyronitrile (AIBN) (e.g., DuPont's Vazo-64^R), 2,2'-azobis (2-amidopropane) hydrochloride (Wako's V-50^R), 2-*t*-butylazo-2-cyanopropane. Most preferred initiators are low temperature azo initiators, such as 2,2'-azobis (2,4-dimethyl-4-methoxyvaleronitrile) (Dupont's Vazo-33^R). The use of low temperature initiation and polymerization leads to higher polymer molecular weights. The concentration of the free radical initiator is preferably 0.001 to 2.0 grams of free radical initiator per 100 grams of total
20 monomer, more preferably 0.01 to 1.0, and most preferably 0.05 to 0.1.

DESCRIPTION OF THE PREFERRED EMBODIMENTS

25 The following Examples no-limitatively illustrate the present invention.

The following nomenclature will be used in the Examples; NVP-RAM = NVP/long chain N-alkylacrylamide/acrylamide terpolymer; NVP-AM = NVP/acrylamide copolymer; PAM = polyacrylamide; SDS = sodium dodecyl sulfate.

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Example 1

Preparation of NVP-RAM Terpolymers (11044-63)

35 A 1,000 ml resin flask was fitted with a condenser, thermometer, stirrer (electric), and nitro-gen inlet and outlet. N-octylacrylamide (0.65 g), NVP (11.88 g) and acrylamide (17.47 g) in 470 g H₂O were polymerized with 0.123 g AIBN in the presence of 1.7 g sodium dodecyl sulfate (SDS) at 45°C for 18 hours. This corresponds to a $[M]/[I]^{1/2} = 60$ (monomer to initiator ratio) and total solids content of 8% in H₂O. Although NVP helps solubilize octylacrylamides, the SDS further homogenizes and clarifies the reaction mixture. The
40 resulting polymer solution was a soluble gel. It was passed through a meat grinder and diluted further with water to 1% concentration. The polymer solution was precipitated in 6 liters of acetone. Copolymers with various ratios of C₈ hydrophobe were prepared and their viscometrics are compared in Table I. The viscosification efficiency of the instant copolymers compared with the composition containing no hydrophobe (NVP/AM/C₈AM = 30/70/0) is clearly evident.

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TABLE I

N-VINYL-PYRROLIDONE-ACRYLAMIDE-N-
OCTYLACRYLAMIDE COPOLYMERS

Polymer Type NVP/AM/C ₈ AM (a)	.15% Polymer, 3.0% NaCl, 0.3% CaCl ₂ Brine Solution, CPS
30/70/0	3.1 (b)
30/69.5/0.5	5.3
30/69.25/0.75	6.8
30/69.0/1.0	17.5

(a) $[M]/[I]^{1/2} = 60$

(b) $[\eta] = 7.02 \text{ dl/g in } 2\% \text{ NaCl}$

Example 2(13006-59P, 60P,70) Effect of SDS in Terpolymerization

The beneficial effect of SDS in the polymerization was demonstrated by preparing terpolymers of NVP/AM/C₈AM with and without 1% SDS. The charged monomer compositions were NVP/AM/C₈AM = 60/39/1. Polymerizations were run at a $[M]/[I]^{1/2} = 60$; total solids in aqueous solution were 18%; polymerizations were run at about 20°C, using Vazo-3^R initiator. The batch with the SDS was clearer during the polymerization than the batch without the SDS, indicating homogenation of the C₈AM monomer. The products were precipitated from acetone and dried. Solution viscosities of the two polymers in 3.0% NaCl and 0.3% CaCl₂ brine were compared. The polymer using SDS had a viscosity of 9.1 cP at 1.28s⁻¹, while the polymer prepared without SDS had a viscosity of only 3.2 cP at 1.28 s⁻¹.

Example 3

The rate of base (OH⁻) catalyzed hydrolysis of NVP-RAM terpolymers compared to NVP/AM copolymers and PAM (polyacrylamide homopolymer) at 40°C is shown in Table II. Surprisingly, the NVP-RAM polymers with varying amounts of C₈ (.5-1% C₈) show less hydrolysis than both RAM and PAM polymers known in the art (Table II).

TABLE II

% HYDROLYSIS VERSUS TIME FOR TERPOLYMERS

Polymer Type	Hydrolysis Time, Hours(a)	% Hydrolysis
PAM	3	22.0
	6	22.5
	12	24.0
RAM (1% C ₈ AM)	3	14.0
	6	15.5
	12	17.0
RAM-NVP (1% C ₈ AM)	3	6.9
	6	6.4
	12	7.4

(a) 40°C, ratio of moles of NaOH/moles polymer = 1/3.

Comparative ExamplePreparation of NVP/AM/N-C₁₀ AM Terpolymers

For comparison of polymers of the present invention with known polymers disclosed in US-A-4,395,524, the following Example was conducted to duplicate the 65AM/30NVP/5N-C₁₀AM (weight percent) polymer of Table VIII therein. This Example corresponds to a 75AM/23NVP/2 mole percent polymer, a $[M]/[I]^{\frac{1}{2}} = 34.0$ and a polymerization solids level of 10%.

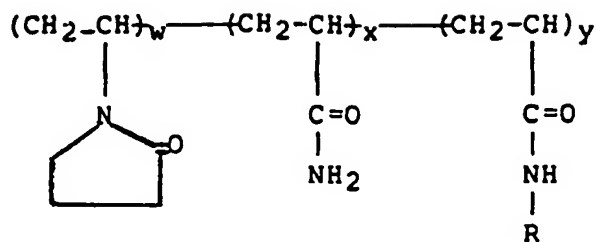
A 250 ml flask is equipped with a thermometer, condenser, air stir motor and an argon inlet tube. Charge reactor with 15.0 g (.135 M) 1-vinyl-2-pyrrolidone, 2.94 g (.041 M) acrylamide, 0.76 g (3.6×10^{-3} M) isomeric N-(n-decyl) acrylamide, 38.7 g tert butanol, 124.5 g deaerated H₂O. Stir and heat mixture to 50°C while maintaining a subsurface argon surge. Add 0.38 g ammonium persulfate in 1.7 g H₂O, 3.4 g isopropanol. Increase reaction temperature to 80° and maintain in argon for 4 hours. Add 0.19 g ammonium persulfate and maintain at 80°C for 1 hour. Evaporate tert butanol on Rotovap, dilute cement with 100 ml methanol and precipitate into acetone. Soak precipitated solid in acetone over night, isolate solid and vacuum dry. Yield is 29.4%. This polymer had a viscosity (.15% polymer, 3.0% NaCl, 0.30% CaCl₂) of only 0.91 cP at 1.28s⁻¹ and a viscosity (.15% polymer, 3.0% NaCl, 0.30% CaCl₂) of 0.94 cP at 11.0s⁻¹. This is clearly a product of much lower viscosification power than those produced by Examples 1 and 2 of the present invention.

Claims

1. A terpolymer having the formula:

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wherein w is 1 to 80 mole percent; x is 10 to 90 mole percent; y is 0.1 to 10.0 mole percent; and R is an alkyl group having 6 to 30 carbon atoms.

2. A terpolymer as claimed in claim 1, wherein w is 10 to 70 mole percent.

3. A terpolymer as claimed in claim 1 or claim 2, wherein x is 30 to 70 mole percent.

4. A terpolymer as claimed in any preceding claim, wherein y is 0.2 to 3.0 per mole percent.

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5. A terpolymer as claimed in any preceding claim, wherein R is an alkyl group of 8 to 18 carbon atoms.

6. The use as a viscosification agent of a terpolymer claimed in any preceding claim.

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